In the Claims:

This listing of claims will replace all prior versions and listings of claims in this application.

1 (withdrawn). An isolated nucleic acid sequence, which encodes a polypeptide with neuronal tryptophane hydroxylase activity, selected from the group of:

- a) a nucleic acid sequence with the sequence depicted in SEQ ID No: 1,
- a nucleic acid sequence that can be deduced from a nucleic acid sequence depicted in SEQ
 ID No: 1 as a consequence of the degenerated genetic code,
- c) derivatives of the nucleic acid sequences depicted in SEQ ID No: 1, which encode polypeptides according to SEQ ID No: 2, which display at least 80% homology at the amino acid level, wherein the biological activity of the polypeptides is not reduced, and
- human genomic nucleic acid sequences, which contain the gene for sn-TPH and exhibit polymorphisms.

2 (eurrently amended). [[The]]An isolated polypeptide encoded by a nucleic acid sequence according to claim 1, which encodes a polypeptide with neuronal tryptophane hydroxylase activity, wherein the nucleic acid sequence is selected from the group of:

- a) a nucleic acid sequence with the sequence depicted in SEQ ID No: 1,
- a nucleic acid sequence that can be deduced from a nucleic acid sequence depicted in SEQ
 ID No: 1 as a consequence of the degenerated genetic code,
- c) derivatives of the nucleic acid sequences depicted in SEQ ID No: 1, which encode polypeptides according to SEQ ID No: 2, which display at least 90% homology at the amino acid level, wherein the tryptophan hydroxylase activity of the polypeptides is not reduced, and
- d) human genomic nucleic acid sequences, which contain the gene for sn-TPH and exhibit polymorphisms.

- 3 (previously presented). The polypeptide according to claim 2, having the sequence depicted in SEO ID No: 2.
- 4 (withdrawn). A recombinant nucleic acid molecule comprising a nucleic acid sequence according to claim 1 or parts of this nucleic acid sequence, wherein the nucleic acid sequence is connected in an anti-sense or sense-direction with one or several regulatory sequences.
 - 5 (withdrawn). A vector comprising a nucleic acid sequence according to claim 1.
- 6 (withdrawn). A recombinant prokaryotic or eukaryotic host organism containing at least one nucleic acid sequence according to claim 1.
- 7 (withdrawn). The recombinant prokaryotic or cukaryotic host organism according to claim 6, wherein the organism is a microorganism or an animal.
- 8 (withdrawn). Use of a polypeptide according to claim 2 or of peptide fragments thereof as an antigen for the production of specific polyclonal or monoclonal antibodies or antibody mixtures.
- 9 (withdrawn). A polyclonal or monoclonal antibody or antibody mixtures, which recognises specific polypeptides according to claim 2.
- 10 (withdrawn). A method for isolating a compound that binds to a polypeptide of claim 2, or of producing a pharmaceutical composition, comprising:
- (a) contacting a mammalian cell that expresses the polypeptide of claim 2 having sn-TPH activity with a compound;
- (b) detecting the presence of a compound that binds to the sn-TPH polypcptide, and
- (c) determining whether the compound binds said sn-TPH polypeptide.

11 (withdrawn). The method, according to claim 10, for the production of a pharmaccutical composition comprising the steps of the process according to claim 10 and the subsequent step of formulating the compound identified in step (e) and/or its pharmaccutically acceptable salts in a pharmaccutically acceptable form.

12 (withdrawn). A method for the treatment of a neuronal disease in a patient, wherein said method is characterised in that serotonin production in the patient is increased or decreased by affecting the snTPH-activity.

13 (withdrawn). The method for the treatment of a neuronal disease according to claim 12, characterised in that the serotonin production is increased by a tissue-specific overexpression of snTPH, by the addition of the precursor substance 5-hydroxy-tryptophane or by the addition of substituted analogues of 5-hydroxy-tryptophane.

14 (withdrawn). The method for the treatment of a neuronal disease according to claim 12, characterised in that the serotonin production is decreased by ribozymes, by antisense-oligonucleotides, by antisense-RNA-expression or by means of a specific TPH-inhibitor.

15 (withdrawn). A method for determining the pharmacogenetic properties of a pharmaceutically active compound and/or improving treatment of a disease, comprising a) administering the compound to a mammal, b) determining the level of expression of snTPH in a biological sample obtained from said mammal, and c) comparing said level of expression of snTPH with a level obtained from a control sample.

16 (withdrawn). The method, according to claim 15, for the improved treatment of a disease, comprising performing the method of claim 15, and increasing or decreasing the doses of the pharmaceutically active compound to be applied to said patient.

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17 (withdrawn). The method according to claim 16, wherein the disease is selected from the group consisting of neuronal diseases, sleep disturbances, anxiety, alcoholism, drug abuse, disorders of food uptake and sexual disorders.

18 (withdrawn). A method for the treatment of sleep disturbances, anxiety, alcoholism, drug abuse, disorders of food uptake or sexual disorders, characterised in that the serotonin level is affected by modulating the gene expression of snTPH wherein said method comprises administering, to a patient in need of such treatment, a nucleotide sequence of claim 1 or a polypeptide of claim 2.

19 (withdrawn). A method for diagnosing a neuronal disease, characterised in that a specific inhibition of the peripheral serotonin biosynthesis is accomplished, followed by subsequently detecting the metabolite concentrations stemming from the CNS and by determining the severity of the disease via a comparative graph.

20 (withdrawn). Use of a nucleic acid sequence according to claim 1 or of a polypeptide according to claim 2 wherein said use is selected from the group consisting of: identifying/discovering proteins, which have specific binding affinities for a polypeptide according to claim 2 or for identifying nucleic acids, which encode for proteins having specific binding affinities for a polypeptide according to claim 2; the isolation of a genomic sequence by means of homology screening or as a marker for human hereditary diseasees; and gene therapy.

21 (withdrawn). The method, according to claim 20, characterised in that the Two-Hybrid-System is employed.

22 - 23 (Canceled).

24 (withdrawn). Use of a DNA sequence according to claim 1 or of a polypeptide according to claim 2 for affecting the serotonin level via specific regulation of the snTPH-activity/amount.

25 (currently amended). A combination therapeutic comprising a polypeptide according to claim 2 and at least one additional protein, in particular for the regulation of the serotonin metabolism.

26 (previously presented). The combination therapeutic according to claim 25, characterised in that the additional protein is a peripheral tryptophane hydroxylase.

27 (cancelled).

28 (withdrawn). Use of the combination therapeutic according to claim 25 for the treatment of bleeding episodes in the psycho-pharmacological treatment of depressions with antidepressants, which affect the serotonin reuptake-transporter, containing antidepressants and von Willebrandfactor.